

Amendments to the claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (original) A method of analyzing the spatial distribution of at least one chemical substance retained by a biological matter, characterized by the steps of
 - (a) supplying a sample of said biological matter as a specimen surface;
 - (b) producing at least one imprint of said specimen surface on at least one corresponding separate substrate surface, said at least one chemical substance being transferred to the same with retained lateral distribution thereon;
 - (c) subjecting said at least one imprint to imaging mass spectrometry, at least one signal from at least two points being produced, the magnitude of said at least one signal being dependent on the amount of said at least one chemical substance laterally present on said substrate surface;
 - (d) recording said at least one signal from said at least two points; and
 - (e) determining said spatial distribution of said at least one chemical substance from said at least one image of said at least one imprint.
2. (original) The method as in claim 1, wherein said at least one chemical substance mainly comprises organic material.
3. (original) The method as in claim 2, wherein said organic material comprises a lipid, an amino acid, a peptide, a protein, a carbohydrate, a nucleotide, a transmitter substance, a drug, or a targeting molecule.

4. (currently amended) The method as in ~~claim 3 and 4~~ claim 3,
wherein said nucleotide is a DNA-molecule.

5. (original) The method as in claim 3, wherein said
targeting molecule is a complementary DNA-sequence.

6. (currently amended) The method as in ~~any of claims 1-3~~ claim 1,
wherein said targeting molecule is an antibody or a fragment thereof.

7. (original) The method as in claim 3, wherein said
targeting molecule comprises a chemical label.

8. (original) The method as in claim 7, wherein said
chemical label is an unusual element or an isotope.

9. (currently amended) The method as in ~~any of claims 1-8~~ claim 1,
wherein said biological matter comprises cells, tissue, virus, body liquid, or biological
molecules.

10. (currently amended) The method as in ~~any of claims 1-9~~ claim 1,
wherein said sample of said biological matter is supplied as a specimen surface *in situ*.

11. (currently amended) The method as in ~~any of claims 1-9~~ claim 1,
wherein said sample of said biological matter is supplied as a specimen surface by
applying it on a solid surface.

12. (original) The method as in claim 11, wherein said solid
surface is a glass surface.

13. (currently amended) The method as in ~~any of claims 1-12~~ claim 1,
wherein multiple sequential imprints are produced from the same area of said specimen
surface.

14. (currently amended) The method as in ~~any of claims 1-13~~ claim 1,
wherein said biological matter is fractured or cut in order to expose its interior before
producing said at least one imprint.

15. (currently amended) The method as in ~~any of claims 1-14~~ claim 1,
wherein said specimen surface is pretreated immediately before producing said at least
one imprint.

16. (original) The method as in claim 15, wherein said
specimen surface is pretreated by condensing a liquid of a non-polar solvent and/or a
polar solvent onto the same.

17. (original) The method as in claim 16, wherein said polar
solvent is a water solution.

18. (currently amended) The method as in ~~claim 16 or 17~~ claim 16,
wherein said specimen surface is first brought to room temperature or cooled and is then
arranged above a heated container containing said liquid.

19. (currently amended) The method as in ~~any of claims 1-18~~ claim 1,
wherein said at least one imprint is produced within 100 s after said pretreatment of said
specimen surface.

20. (currently amended) The method as in ~~any of claims 1-19~~ claim 1,
wherein said specimen and/or said substrate is flexible.

21. (currently amended) The method as in ~~any of claims 1-20~~ claim 1,
wherein said substrate surface is a metal surface.

22. (original) The method as in claim 21, wherein said metal
is silver, gold, palladium, platinum, nickel, chromium, or copper, preferably silver.

23. (currently amended) The method as in ~~any of claims 1-22~~ claim 1,
wherein said substrate surface is structured.

24. (original) The method as in claim 23, wherein said
substrate surface is structured with protrusions of 0.01-5 μ m.

25. (currently amended) The method as in ~~any of claims 1-22~~ claim 1,
wherein said substrate surface is polished.

26. (currently amended) The method as in ~~any of claims 1-25~~ claim 1,
wherein said substrate surface is cleaned immediately before producing said at least one
imprint.

27. (original) The method as in claim 26, wherein said
substrate surface is cleaned by means of chemical etching, plasma cleaning, or UV/ozone
treatment, or a combination thereof.

28. (currently amended) The method as in ~~any of claims 1-27~~ claim 1,
wherein said specimen surface is subjected to lyophilization, freeze-substitution, or air
drying before producing said at least one imprint.

29. (currently amended) The method as in ~~any of claims 1-29~~ claim 1,
wherein said biological matter is subjected to a salt solution before and/or after supplying
said sample of biological matter as a specimen surface.

30. (original) The method as is claim 29, wherein said salt is a
sodium salt, a potassium salt, a copper salt or a silver salt, preferably a silver salt.

31. (currently amended) The method as in ~~any of claims 1-30~~ claim 1,
wherein said at least one imprint is produced by pressing said specimen surface against
said substrate surface.

32. (original) The method as in claim 31, wherein said pressing is accomplished by means of a compressible material.

33. (currently amended) The method as in claim 31 ~~or~~ 32, wherein said pressing is accomplished by applying a force between 0.01 and 10 MPa.

34. (currently amended) The method as in ~~any of claims 31-33~~ claim 31, wherein said pressing is performed for up to 100 s.

35. (currently amended) The method as in ~~any of claims 31-34~~ claim 31, wherein said pressing is performed so that said at least one imprint represents below 5 monolayers, preferably below 2 monolayers, comprising said at least one chemical substance on said substrate surface.

36. (currently amended) The method as in ~~any of claims 21-27~~ claim 21, wherein a metal layer is deposited onto said substrate surface before producing said at least one imprint.

37. (currently amended) The method as in ~~any of claims 1-35~~ claim 1, wherein a metal layer is deposited onto said substrate surface after producing said at least one imprint.

38. (original) The method as in claim 37, wherein said layer of metal has a thickness of less than 100 nm.

39. (currently amended) The method as in ~~any of claims 36-38~~ claim 36, wherein said layer of metal is a silver layer.

40. (currently amended) The method as in ~~any of claims 1-39~~ claim 1, wherein said imaging mass spectrometry is a Secondary Ion Mass Spectrometry.

41. (original) The method as in claim 40, wherein said Secondary Ion Mass Spectrometry is Time of Flight - Secondary Ion Mass Spectrometry.

42. (currently amended) The method as in claim 40 or 41, wherein a focused beam of ions is produced by the primary ion source in said Secondary Ion Mass Spectrometry.

43. (original) The method as in claim 42, wherein said ions are C₆₀, Ga, In, or Au ions.

44. (original) The method as in claim 43, wherein said Au ions are clusters of n ions, n ≤ 10.

45. (original) The method as in claim 42, wherein said focused beam has a diameter below 10 μm, preferably below 1 μm.

46. (currently amended) The method as in ~~any of claims 1-45~~ claim 1, wherein a light sensitive matrix is applied onto said substrate surface before producing said at least one imprint.

47. (currently amended) The method as in ~~any of claims 1-45~~ claim 1, wherein a light sensitive matrix is applied onto said substrate surface after producing said at least one imprint.

48. (currently amended) The method as in ~~any of claims 1-45~~ claim 1, wherein a light sensitive matrix is applied onto said specimen surface before producing said at least one imprint, a portion of said light sensitive matrix being transferred to the substrate surface when said at least one imprint is produced.

49. (currently amended) The method as in ~~any of claims 1-35 and 46-48~~
claim 1, wherein said imaging mass spectrometry is a Matrix Assisted Laser Desorption
Ionisation.

50. (original) The method as in claim 49, wherein the light
source of said Matrix Assisted Laser Desorption Ionization comprises a focused laser
beam, preferably an ultraviolet laser beam.

51. (currently amended) The method as in ~~any of claims 1-50~~ claim 1,
wherein said at least one signal is recorded from an array of points on said substrate
surface.

52. (currently amended) The method as in ~~any of claims 1-51~~ claim 1,
wherein said at least one image is produced from said at least one signal, the colour or the
brightness in each point of said at least one image being dependent on the magnitude of
said at least one signal from the corresponding point on said substrate surface.